Amendments to the Claims.

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

- 1. (Currently amended) A method of producing a radiolabelled gallium complex in a form suitable for use in PET or SPECT radiopharmaceutical imaging, said method comprising by reacting a Ga³⁺ radioisotope in a suitable solvent with a macrocyclic bifunctional chelating agent, wherein said macrocyclic bifunctional chelating agent is linked to a targeting vector selected from the group consisting of proteins, glycoproteins, lipoproteins, polypeptides, glycopolypeptides, lipopolypeptides, peptides, glycopeptides, lipopeptides, carbohydrates, nucleic acids, oligonucleotides or small organic molecules; characterised in that the reaction is carried out using microwave activation at 80 to 120 W for 20 s to 2 min.
- 2. (Previously presented) The method according to claim 1 wherein the Ga³⁺ radioisotope is selected from the group consisting of ⁶⁶Ga³⁺, ⁶⁷Ga³⁺ and ⁶⁸Ga³⁺.
- 3. (Previously presented) The method according to claim 1 wherein the Ga³⁺ radioisotope is 68 Ga $^{3+}$.
- 4. (Cancelled)
- 5. (Previously presented) The method according to claim 1 wherein the macrocyclic bifunctional chelating agent comprises hard donor atoms, preferably O and N atoms.
- 6. (Cancelled)

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7. (Cancelled)

8. (Previously presented) The method according to claim 1 wherein the target vector is a

peptide or oligonucleotide.

9. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out at 90 to 110 W.

10. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out for 30 s to 90 s.

11. (Previously presented) The method according to claim 3 wherein the ⁶⁸Ga³⁺ is obtained

by contacting the eluate from a ⁶⁸Ge/⁶⁸Ga generator with an anion exchanger and eluting

⁶⁸Ga³⁺ from said anion exchanger.

12. (Previously presented) The method according to claim 11 wherein the ⁶⁸Ge/⁶⁸Ga

generator comprises a column comprising titanium dioxide.

13. (Previously presented) The method according to claim 11 wherein the anion exchanger

comprises HCO₃ as counterions.

14. (Previously presented) The method according to claim 11 wherein the anion exchanger is

an anion exchanger comprising quaternary amine functional groups, or the ion exchanger

is a anion exchange resin based on polystyrene-divinylbenzene.

15. (Previously presented) The method according to claim 1 for the production of ⁶⁸Ga-

radiolabelled PET tracers.

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16. (Withdrawn) Method according to claim 11 wherein the eluting ⁶⁸Ga³⁺ is in the picomolar to nanomolar range after the elution, and more preferably in a nanomolar to micromolar level.